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Treatment of central retinal vein occlusion by injection of
tissue plasminogen activator into a retinal vein .
Weiss, Jeffrey N.
American Journal of Ophthalmology, v126, n1, p142(3)
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1998
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PURPOSE: To report the injection of tissue plasminogen
activator

into a retinal vein to treat central retinal vein occlusion .

METHODS: An 81-year-old woman with visual loss of the right eye
secondary to central retinal vein occlusion developed central
retinal vein occlusion and visual loss in her left eye.

Treatment of
her left eye with topical ocular hypotensive medications,
pentoxifylline,
and laser chorioretinal anastomosis was without benefit. Thereafter,
she
underwent vitreoretinal surgery, including tissue plasminogen
activator
injection into a branch retinal vein of her left eye.

RESULTS: The patient reported subjective improvement in the
vision of
her left eye. Ophthalmoscopic and fluorescein angiographic improvement
were
also noted.

CONCLUSION: The feasibility of cannulating a retinal vein for
treatment has been demonstrated. (Am J Ophthalmol 1998;126:142-144.
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CENTRAL RETINAL VEIN OCCLUSION IS A COMMON retinal -
vascular

disorder with potentially blinding complications. An increased risk of
central retinal vein occlusion has been found in patients with
systemic hypertension, diabetes mellitus, and open-angle glaucoma.(1)
Histopathologic studies have demonstrated thrombosis at the lamina
cribrosa.(2) The anatomy of the lamina cribrosa provide a mechanical
factor

for thrombosis and an increase in coagulability, increase in blood
viscosity, and/or venous stasis provide a hematologic predisposition
for
thrombosis.(3)

An 80-year-old woman with hypercholesterolemia experienced a

central
retinal vein occlusion of the right eye. One month later,
rubeosis
irides and neovascular glaucoma were noted, and panretinal laser
photocoagulation was performed. The right eye was stable 1 year later,
with
a best-corrected visual acuity of RE, 8/200, when she presented with
central retinal vein occlusion in her left eye. Treatment with
topical ocular hypotensive medications and pentoxifylline were without
benefit. A laser chorioretinal anastomosis was attempted but was not
successful, and her best-corrected visual acuity declined to 20/400,
with
extensive hemorrhages observed in all retinal quadrants.

After extensive discussion and mandatory second opinion, the
patient
agreed to undergo experimental retinal surgery for this condition.
Informed consent included the risk of retinal detachment and
intraocular
hemorrhage resulting from cannulating the retinal vein and tissue
plasminogen activator injection. The procedure was approved by the
Institutional Review Board.

On the left eye, a standard three-port vitrectomy was initially
performed. Using a 33-gauge needle and mechanical micromanipulator
that I
had designed, an additional sclerotomy was made so that the needle
would be
parallel to the lumen of the selected superior branch retinal vein
near
the optic disk. A chandelier light source was used to minimize
intraocular
movement. The patient's intraocular pressure was lowered to 5 mm Hg,
and
spontaneous venous pulsations were observed. The vessel was entered by
remote mechanical control, which maintained ocular stability, and
there was
no hemorrhaging. A bolus of 20 (micro)g/0.1 ml of tissue plasminogen
activator was injected toward the optic nerve head. Intraocular
pressure
was raised to 25 mm Hg, and the cannula was removed without
hemorrhaging
from the retinal vessel. When her intraocular pressure was raised to
30
mm Hg, venous pulsations were produced. Laser photocoagulation or gas
tamponade was not required. There were no intraoperative or
postsurgical
complications.

A preoperative fluorescein angiogram of the left eye
demonstrated
venous filling at 33 seconds, whereas the same study performed
approximately 3 weeks postoperatively showed venous filling at 24
seconds.
Subsequent to the surgery, there was a notable decrease in the amount

of

retinal hemorrhaging observed, especially nasally.

Approximately 3 months postoperatively, and though the patient reported an improvement in vision, neovascularization of the angle was observed. Panretinal laser photocoagulation was performed with complete

resolution of the neovascularization and an improvement in best-corrected

visual acuity to LE, 20/400. Approximately 9 months postvitrectomy surgery and 5 months postlaser photocoagulation, the best-corrected visual

acuity was stable at LE, 20/400, and the patient remained pleased with her

perceived visual improvement.

The cannulation of retinal vessels in animals has been described,(4) as has tissue plasminogen activator therapy for central

retinal vein occlusion in patients. To the best of my knowledge, this

is the first report of cannulation of a retinal vein with injection

of tissue plasminogen activator in a patient with a central retinal vein occlusion.(5)

This case demonstrates that it is clinically possible to cannulate a

retinal vessel and safely infuse a drug. I am presently developing a smaller cannula and improving the ease of use of the micromanipulator. Whether this technique will prove beneficial in the treatment of central

retinal vein occlusion, ocular tumors, and other retinal and retinal -vascular disorders that would benefit from local targeting of drug

treatment remains to be determined by larger controlled clinical studies.

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